

Seat No.: _____

Enrolment No. _____

GUJARAT TECHNOLOGICAL UNIVERSITY
B.PHARM – SEMESTER – 6 EXAMINATION – SUMMER-2025

Subject Code: BP604TT

Date: 17-05-2025

Subject Name: Biopharmaceutics and Pharmacokinetics

Time: 10.30 AM TO 01.30 PM

Total Marks: 80

Instructions:

1. Attempt any five questions.
2. Make suitable assumptions wherever necessary.
3. Figures to the right indicate full marks.

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|-------------|---|-----------|
| Q.1 | (a) Discuss the physicochemical factors affecting the GI absorption of drugs. | 06 |
| | (b) Explain briefly about different mechanism of drug absorption. | 05 |
| | (c) Define drug distribution. Write a short note on volume of distribution. | 05 |
| Q.2 | (a) Define clearance and write about renal clearance in detail. | 06 |
| | (b) Write a short note on kinetics of protein-drug binding. | 05 |
| | (c) Discuss factors affecting renal excretion of drugs. | 05 |
| Q.3 | (a) Differentiate absolute and relative bioavailability. Discuss the pharmacokinetic methods for the bioavailability measurement. | 06 |
| | (b) Write a note on IVIVC. | 05 |
| | (c) Differentiate between plasma-protein drug binding and tissue-drug binding. | 05 |
| Q.4 | (a) What are pharmacokinetic models? What are applications of such models? Discuss any one pharmacokinetic model. | 06 |
| | (b) What do you mean by nonlinear pharmacokinetics? Discuss factors causing Nonlinearity. | 05 |
| | (c) Explain Michaelis-menton equation. | 05 |
| Q.5 | (a) Discuss briefly the concept of loading dose and maintenance dose. | 06 |
| | (b) Write a brief note on two compartment open model. | 05 |
| | (c) Explain merits and demerits of non-compartmental pharmacokinetics. | 05 |
| Q. 6 | (a) Write a brief note on physiological models. | 06 |
| | (b) Write on methods to enhance dissolution rate of poorly soluble drugs. | 05 |
| | (c) Derive equation in determination of absorption rate constant (Ka) using Wegner Nelson Method. | 05 |
| Q.7 | (a) List out the various pharmacokinetic parameters and define. | 06 |
| | (b) Describe Latin Square cross over design for bioequivalence study. | 05 |
| | (c) Explain physiological barriers to drug distribution in body. | 05 |
